=> s vascular endothelial growth factor

15734 VASCULAR

4014 ENDOTHELIAL

125665 GROWTH

231766 FACTOR

L1 67 VASCULAR ENDOTHELIAL GROWTH FACTOR
(VASCULAR (W) ENDOTHELIAL (W) GROWTH (W) FACTOR)

=> s vegf3

L2 0 VEGF3

=> s veqf

L3 42 VEGF

=> s 11 or 13

L4 72 L1 OR L3

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L4: 6 of 72 US PAT NO: 5,654,404 : IMAGE AVAILABLE:

Aug. 5, 1997 DATE ISSUED:

Protection against liver damage by HGF TITLE:

Filip Roos, Brisbane, CA INVENTOR: Ralph Schwall, Pacifica, CA

Genentech, Inc., So. San Francisco, CA (U.S. corp.) ASSIGNEE:

08/419,654 APPL-NO: DATE FILED: Apr. 10, 1995

Division of Ser. No. 310,361, Sep. 21, 1994, which is a REL-US-DATA: continuation of Ser. No. 968,711, Oct. 30, 1992,

abandoned, which is a continuation-in-part of Ser. No.

946,263, Sep. 16, 1992, abandoned.

:6: C07; 14/435; C12P 21/08; A61K 39/00; A61K 38/16 530/38 , 350; 424/134.1, 136.1, 178.1 INT-CL:

US-CL-ISSUED: US-CL-CURRENT: 530/387.3; 424/134.1, 136.1, 178.1; 530/550

530/389.2, 387.3, 399, 350; 424/138.1, 124.1, 145.1; SEARCH-FLD:

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186 ART-UNIT:

Lila Feisee PRIM-EXMR: ASST-EXMR: John Lucas

Merchant, Gould, Smith, Edell, Welter & Schmidt LEGAL-REP:

ABSTRACT:

The present invention provides methods for preventing occurrence or progression of liver damage using hepatocyte growth factor. In the methods, a preventatively effective amount of the hepatocyte growth factor is administered to the patient. The hepatocyte growth factor can be administered, for instance, prior to administering a hepatotoxic therapy to the patient. The hepatocyte growth factor can further be administered with activin or transforming growth factor-beta to prevent liver damage. Compositions comprising hepatocyte growth factor and activin antagonist or transforming growth factor-beta antagonist are also provided by the invention.

18 Claims, 9 Drawing Figures

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- 40. 5,506,107, Apr. 9, 1996, Selecting ligand agonists and antagonists; Brian C. Cunningham, et al., 435/7.21, 7.8; 436/501, 537 :IMAGE AVAILABLE:

=> d 30 clms

US PAT NO: 5,607,918 : IMAGE AVAILABLE: L4: 30 of 72

CLAIMS:

CLMS(1)

What is claimed is:

1. An isolated protein having the property of promoting proliferation of endothelial cells or mesodermal cells, said isolated protein comprising a sequence of amino acids selected from the group consisting of the amino

acid sequence of FIG. (SEQ ID NO:2), the amino acid sequence of FIG. 2 (SEQ ID NO:3), the am acid sequence of FIG. 4 (SEQ NO:5), the amino acid sequence of FIG. 6 (SEQ ID NO:7), the amino acid sequence of FIG. 8 (SEQ ID NO:9), and the amino acid sequence of FIG. 11 (SEQ ID NO:11).

CLMS(2)

2. An isolated protein according to claim 1, wherein said protein comprises the amino acid sequence of FIG. 1 (SEQ ID NO:2).

CLMS(3)

3. An isolated protein according to claim 1, wherein said isolated protein comprises the amino acid sequence of FIG. 2 (SEQ ID NO:3).

CLMS (4)

4. An isolated protein according to claim 1, wherein said isolated protein comprises the amino acid sequence of FIG. 4 (SEQ ID NO:5).

CLMS(5)

5. An isolated protein according to claim 1, wherein said isolated protein comprises the amino acid sequence of FIG. 6 (SEQ ID NO:7).

CLMS(6)

6. An isolated protein according to claim 1, wherein said isolated protein comprises the amino acid sequence of FIG. 8 (SEQ ID NO:9).

CLMS(7)

7. An isolated protein according to claim 1, wherein said isolated protein comprises the amino acid sequence of FIG. 11 (SEQ ID NO:11).

CLMS(8)

8. An isolated protein according to claim 1, wherein said isolated protein is a mammalian protein.

CLMS(9)

9. An isolated protein according to claim 8, wherein said mammalian protein is a murine protein.

CLMS (10)

10. An isolated protein according to claim 8, wherein said mammalian protein is a human protein.

CLMS (11)

11. An isolated protein according to claim 1, wherein said isolated protein promotes proliferation of vascular endothelial cells.

CLMS (12)

12. An isolated protein produced by expression of a DNA selected from the group consisting of the DNA of FIGS. 1 and 2 (SEQ ID NO:1), the DNA of FIG. 3 (SEQ ID NO:4), the DNA of FIG. 5 (SEQ ID NO:6), the DNA of FIG. 7 (SEQ ID NO:8), the DNA of FIG. 10 (SEQ ID NO:10), and DNA which hybridizes under stringent conditions with at least one of the foregoing DNA sequences.

CLMS (13)

13. A pharmaceutical emposition comprising an effect endothelial or mesodermal cell proliferation promoting amount of an isolated protein according to claim 1, and at least one pharmaceutical carrier or diluent.

=> d 41-60

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- 45. 5,470,878, Nov. 28, 1995, Cell signaling inhibitors; John Michnick, et al., 514/558, 258, 262, 274, 299, 315, 418, 425, 529, 552, 561, 613, 617, 626, 629, 669; 544/254, 285, 301; 546/183, 243; 548/486, 556 :IMAGE AVAILABLE:
- 46. 5,464,815, Nov. 7, 1995, Inhibition of heparin-binding; Steven Chamow, et al., 514/8; 424/85.2; 436/86, 87; 514/21; 530/412 :IMAGE AVAILABLE:
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- 51. 5,407,810, Apr. 18, 1995, Aqueous multiple-phase isolation of polypeptide; Stuart Builder, et al., 435/69.1, 804; 530/399, 412, 422, 808 :IMAGE AVAILABLE:
- 52. 5,391,164, Feb. 21, 1995, Subcutaneous implantable multiple-agent delivery system; Vincent C. Giampapa, 604/891.1; 424/424, 425 :IMAGE AVAILABLE:
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- 54. 5,382,514, Jan. 17, 1995, In vivo angiogenesis assay; Antonino Passaniti, et al., 435/7.21; 424/520, 572; 435/7.23, 29; 436/63, 64, 813 :IMAGE AVAILABLE:

- 55. 5,342,763, Aug. 3 1994, Method for producing polypeptide via bacterial fermentation James R. Swartz, 435/69.1, 71. 530/401 :IMAGE AVAILABLE:
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- 57. 5,332,671, Jul. 26, 1994, Production of vascular endothelial cell growth factor and DNA encoding same; Napoleone Ferrara, et al., 435/360, 69.4, 69.6, 320.1; 536/23.5, 23.51 :IMAGE AVAILABLE:
- 58. 5,329,028, Jul. 12, 1994, Carbohydrate-directed cross-linking reagents; Avi J. Ashkenazi, et al., 548/548, 546, 547, 549 :IMAGE AVAILABLE:
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- 60. 5,326,568, Jul. 5, 1994, Method of tissue-specific delivery; Vincent C. Giampapa, 424/426, 423, 424; 604/890.1, 892.1; 623/11 :IMAGE AVAILABLE:

FILE 'HOME' ENTERED AT 09:55:03 ON 09 SEP 1997

=> s vascular endothelial growth factor or vegf

THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE Some commands only work in certain files. For example, the EXPAND command can only be used to look at the index in a file which has an index. Enter "HELP COMMANDS" at an arrow prompt (=>) for a list of commands which can be used in this file.

=> file medline

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MEDLINE ANNUAL RELOAD AVAILABLE ON STN IN RECORD TIME (2/08/97). ENTER HELP RLOAD FOR DETAILS.

THIS FILE CONTAINS CAS REGISTRY NUMBERS FOR EASY AND ACCURATE SUBSTANCE IDENTIFICATION.

=> s vascular endothelial growth factor or vegf

197586 VASCULAR

47598 ENDOTHELIAL

440125 GROWTH

339460 FACTOR

822 VASCULAR ENDOTHELIAL GROWTH FACTOR

(VASCULAR (W) ENDOTHELIAL (W) GROWTH (W) FACTOR)

707 VEGF

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707 VEGF

1463151 3

2 VEGF-3

(VEGF (W) 3)

L2 2 VEGF3 OR VEGF-3

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L2 ANSWER 1 OF 2 MEDLINE

AN 97342551 MEDLINE

TI Sequencing of the human vascular endothelial growth factor (**VEGF**) 3' untranslated region (UTR): conservation of five hypoxia-inducible RNA-protein binding sites.

AU Levy N S; Goldberg M A; Levy A P

CS Department of Medicine, Georgetown University Medical Center, Washington, DC 20007, USA.

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                        BI)
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SO
     Journal code: AOW. ISSN: 0006-3002.
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     Netherlands
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     Journal; Article; (JOURNAL ARTICLE)
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     GENBANK-Y08736; GENBANK-J03145
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     ANSWER 2 OF 2 MEDLINE
L2
                  MEDLINE
     96162020
ΑN
     Post-transcriptional regulation of vascular endothelial growth
TI
     factor by hypoxia.
ΑU
     Levy A P; Levy N S; Goldberg M A
     Cardiology Division, Brigham and Women's Hospital, Boston,
CS
     Massachusetts 02115, USA.
NC
     T32HL07604 (NHLBI)
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     Journal code: HIV. ISSN: 0021-9258.
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     Journal; Article; (JOURNAL ARTICLE)
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     ANSWER 1 OF 6 MEDLINE
L3
                  MEDLINE
AN
     97332628
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     Enholm B; Paavonen K; Ristimaki A; Kumar V; Gunji Y; Klefstrom J;
ΑU
     Kivinen L; Laiho M; Olofsson B; Joukov V; Eriksson U; Alitalo K
     Molecular/Cancer Biology Laboratory, University of Helsinki,
CS
     Finland.
     ONCOGENE, (1997 May 22) 14 (20) 2475-83.
SO
     Journal code: ONC. ISSN: 0950-9232.
     ENGLAND: United Kingdom
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     Journal; Article; (JOURNAL ARTICLE)
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     English
     Priority Journals; Cancer Journals
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     19970901
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     ANSWER 2 OF 6 MEDLINE
L3
                  MEDLINE
     97167624
ΑN
     Expression of vascular endothelial growth factor and placenta growth
ΤI
     factor in human placenta.
     Vuorela P; Hatva E; Lymboussaki A; Kaipainen A; Joukov V; Persico M
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G; Alitalo K; Halmesmaki E

Department of Objectics and Gynecology, Helsinki piversity Central Hospital, Finland piia.vuorela@helsinki.fi CS BIOLOGY OF REPRODUCTION, (1997 Feb) 56 (2) 489-94. so Journal code: A3W. ISSN: 0006-3363. CY United States DT Journal; Article; (JOURNAL ARTICLE) LΑ English FS Priority Journals EM9706 19970604 EW ANSWER 3 OF 6 MEDLINE L3 97164697 MEDLINE ANVEGF-C receptor binding and pattern of expression with VEGFR-3 ΤI suggests a role in lymphatic vascular development. Kukk E; Lymboussaki A; Taira S; Kaipainen A; Jeltsch M; Joukov V; ΑU Alitalo K Molecular/Cancer Biology Laboratory, Haartman Institute, University CS of Helsinki, Finland. DEVELOPMENT, (1996 Dec) 122 (12) 3829-37. SO Journal code: ECW. ISSN: 0950-1991. ENGLAND: United Kingdom CY Journal; Article; (JOURNAL ARTICLE) DT English LΑ Priority Journals FS GENBANK-X68203; GENBANK-U73620 OS EΜ 9705 19970501 EW ANSWER 4 OF 6 MEDLINE L3 AΝ 96325041 MEDLINE ΤI Genomic organization of the mouse and human genes for vascular endothelial growth factor B (VEGF-B) and characterization of a second splice isoform. Olofsson B; Pajusola K; von Euler G; Chilov D; Alitalo K; Eriksson U ΑU Ludwig Institute for Cancer Research, Stockholm Branch, Box 240, CS S-171 77 Stockholm, Sweden. JOURNAL OF BIOLOGICAL CHEMISTRY, (1996 Aug 9) 271 (32) 19310-7. SO Journal code: HIV. ISSN: 0021-9258. United States CY Journal; Article; (JOURNAL ARTICLE) DT LΑ English FS Priority Journals; Cancer Journals GENBANK-U52819; GENBANK-U52820 os EM9611 ANSWER 5 OF 6 MEDLINE L3MEDLINE 96220114 ΑN Novel human vascular endothelial growth factor genes VEGF-TI B and VEGF-C localize to chromosomes 11q13 and 4q34, respectively. Paavonen K; Horelli-Kuitunen N; Chilov D; Kukk E; Pennanen S; ΑU Kallioniemi O P; Pajusola K; Olofsson B; Eriksson U; Joukov V; Palotie A; Alitalo K Molecular/Cancer Biology Laboratory, Haartman Institute, Helsinki, CS Finland. CIRCULATION, (1996 Mar 15) 93 (6) 1079-82. so Journal code: DAW. ISSN: 0009-7322. CY United States Journal; Article; (JOURNAL ARTICLE) DT

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- L3 ANSWER 6 OF 6 MI INE
- AN 96197355 MEDL-NE
- Vascular endothelial growth factor B, a novel growth factor for endothelial cells.
- AU Olofsson B; Pajusola K; Kaipainen A; von Euler G; Joukov V; Saksela O; Orpana A; Pettersson R F; Alitalo K; Eriksson U
- CS Ludwig Institute for Cancer Research, Stockholm, Sweden.
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- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Cancer Journals; Priority Journals
- OS GENBANK-U48800; GENBANK-U48801
- EM 9609
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- L3 ANSWER 1 OF 6 MEDLINE
- The vascular endothelial growth factor (VEGF) family has recently AB been expanded by the isolation of two additional growth factors, **VEGF-B** and VEGF-C. Here we compare the regulation of steady-state levels of VEGF, VEGF-B and VEGF-C mRNAs in cultured cells by a variety of stimuli implicated in angiogenesis and endothelial cell physiology. Hypoxia, Ras oncoprotein and mutant p53 tumor suppressor, which are potent inducers of VEGF mRNA did not increase VEGF-B or VEGF-C mRNA levels. Serum and its component growth factors, platelet-derived growth factor (PDGF) and epidermal growth factor (EGF) as well as transforming growth factor-beta (TGF-beta) and the tumor promoter phorbol myristate 12,13-acetate (PMA) stimulated VEGF-C, but not VEGF-B mRNA expression. Interestingly, these growth factors and hypoxia simultaneously downregulated the mRNA of another endothelial cell specific ligand, angiopoietin-1. Serum induction of VEGF-C mRNA occurred independently of protein synthesis; with an increase of the mRNA half-life from 3.5 h to 5.5-6 h, whereas **VEGF-B** mRNA was very stable (T 1/2>8 h). Our results reveal that the three VEGF genes are regulated in a strikingly different manner, suggesting that they serve distinct, although perhaps overlapping functions in vivo.
- L3 ANSWER 2 OF 6 MEDLINE
- Normal development and function of the placenta requires invasion of AΒ the maternal decidua by trophoblasts, followed by abundant and organized vascular growth. Little is known of the significance and function of the vascular endothelial growth factor (VEGF) family, which includes VEGF, VEGF-B, and VEGF-C, and of placenta growth factor (PIGF) in these processes. In this study we have analyzed the expression of VEGF and PIGF mRNAs and their protein products in placental tissue obtained from noncomplicated pregnancies. Expression of VEGF and PIGF mRNA was observed by in situ hybridization in the chorionic mesenchyme and villous trophoblasts, respectively. Immunostaining localized the VEGF and PIGF proteins in the vascular endothelium, which was defined by staining for von Willebrand factor and for the Tie receptor tyrosine kinase, an early endothelial cell marker. VEGF-B and VEGF-C mRNAs were strongly expressed in human placenta as evidenced by Northern blot analysis. These data imply that VEGF and PIGF are produced by different cells but that both target the endothelial cells of normal human term placenta.

The vascular end elial growth factor family has cently been expanded by the isolation of two new VEGF-related ctors, VEGF-B and VEGF-C. The physiological functions of these factors are largely unknown. Here we report the cloning and characterization of mouse VEGF-C, which is produced as a disulfide-linked dimer of 415 amino acid residue polypeptides, sharing an 85% identity with the human VEGF-C amino acid sequence. The recombinant mouse VEGF-C protein was secreted from transfected cells as VEGFR-3 (Flt4) binding polypeptides of 30-32x10(3) Mr and 22-23x10(3) Mr which preferentially stimulated the autophosphorylation of VEGFR-3 in comparison with VEGFR-2 (KDR). In in situ hybridization, mouse VEGF-C mRNA expression was detected in mesenchymal cells of postimplantation mouse embryos, particularly in the regions where the lymphatic vessels undergo sprouting from embryonic veins, such as the perimetanephric, axillary and jugular regions. In addition, the developing mesenterium, which is rich in lymphatic vessels, showed strong VEGF-C expression. VEGF-C was also highly expressed in adult mouse lung, heart and kidney, where VEGFR-3 was also prominent. The pattern of expression of VEGF-C in relation to its major receptor VEGFR-3 during the sprouting of the lymphatic endothelium in embryos suggests a paracrine mode of action and that one of the functions of VEGF-C may be in the regulation of angiogenesis of the lymphatic vasculature.

ANSWER 4 OF 6 MEDLINE L3

AΒ

A second isoform and the genomic structures of mouse and human AΒ vascular endothelial growth factor B are described. Both genes consist of seven coding exons and span about 4 kilobases of DNA. The two identified isoforms of vascular endothelial growth factor B are generated by alternative splicing where different splice acceptor sites in exon 6 introduce a frameshift and a partial use of different but overlapping reading frames. Consequently, the COOH-terminal domains in the two isoforms show no resemblance. Mouse and human cDNA clones for the novel isoform of vascular endothelial growth factor B encoded a secreted protein of 186 amino acid residues. Expression in transfected cells generated a protein of 25 kDa which upon secretion was modified by O-linked glycosylation and displayed a molecular mass of 32 kDa under reducing conditions. The protein was expressed as a disulfide-linked homodimer, and heterodimers were generated when coexpressed with vascular endothelial growth factor. The entirely different COOH-terminal domains in the two isoforms of vascular endothelial growth factor B imply that some functional properties of the two proteins are distinct.

ANSWER 5 OF 6 MEDLINE L3

BACKGROUND: Vascular endothelial growth factor (VEGF) is an important regulator of endothelial cell proliferation, migration, and permeability during embryonic vasculogenesis as well as in physiological and pathological angiogenesis. The recently isolated VEGF-B and VEGF-C cDNAs encode novel growth factor genes of the VEGF family. METHODS AND RESULTS: Southern blotting and polymerase chain reaction analysis of somatic cell hybrids and fluorescence in situ hybridization (FISH) of metaphase chromosomes were used to assess the chromosomal localization of VEGF-B and VEGF-C genes. The VEGF-B gene was found on chromosome 11q13, proximal to the cyclin D1 gene, which is amplified in a number of human carcinomas. However, VEGF-B was not amplified in several mammary carcinoma cell lines containing amplified cyclin D1. The VEGF-C gene was located on chromosome 4q34, close to the human aspartylglucosaminidase gene previously mapped to 4q34-35. CONCLUSIONS: The VEGF-B locus in 11q13 and the VEGF-C locus in 4q34 are candidate targets for mutations that lead to vascular malformations or cardiovascular diseases.

L3 ANSWER 6 OF 6 ME. INE

We have isolated and characterized a novel growth factor for endothelial cells, vascular endothelial growth factor B (
VEGF-B), with structural similarities to vascular endothelial growth factor (VEGF) and placenta growth factor.

VEGF-B was particularly abundant in heart and skeletal muscle and was coexpressed with VEGF in these and other tissues. VEGF-B formed cell-surface-associated disulfide-linked homodimers and heterodimerized with VEGF when coexpressed. Conditioned medium from transfected 293EBNA cells expressing VEGF-B stimulated DNA synthesis in endothelial cells. Our results suggest that VEGF-B has a role in angiogenesis and endothelial cell growth, particularly in muscle.